

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 260 545 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:

27.11.2002 Bulletin 2002/48

(51) Int Cl.7: C08K 5/00, C08L 23/10,

F16L 9/12, F16L 9/127,

F16L 9/133, B32B 1/08,

B32B 27/32, C08F 110/06,

C08F 297/08, C08F 10/06

(21) Application number: 01112365.0

(22) Date of filing: 21.05.2001

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR

Designated Extension States:

AL LT LV MK RO SI

(71) Applicant: Borealis Technology OY
06201 Porvoo (FI)

(72) Inventor: The designation of the inventor has not yet been filed

(74) Representative: VA TECH Patente GmbH & Co
Stahlstrasse 21 a
4020 Linz (AT)

(54) Industrial polyolefin piping system

(57) Industrial polyolefin piping system with improved stiffness, impact strength for use at high service temperature, comprising single- or multilayer pipes, fittings, chambers, valves and vessels, consisting of β -nucleated propylene homopolymers with an $IR\tau > 0.98$ having a tensile modulus ≥ 1500 MPa, a Charpy impact strength, notched, at $+23^\circ\text{C} \geq 30$ kJ/m² and a Vicat B temperature $> 90^\circ\text{C}$ and a Heat Distortion Temperature

$> 100^\circ\text{C}$.

The industrial polyolefin piping system is suitable for chemical plant constructions comprising single- or multilayer pipes, fittings, chambers, valves and vessels with improved stiffness, impact strength and high service temperature, preferred for conveyance of natural gas, dangerous liquids and/or toxic liquids.

Description

Field of the invention

- 5 [0001] The invention relates to an industrial polyolefin piping system with improved stiffness, impact strength for use at high service temperature, comprising single or multilayer pipes, fittings, valves, chambers and vessels, consisting of a propylene homopolymer, as well as a process for producing them.

Background of the invention

- 10 [0002] Pipes, injection molded parts and blow molded parts from propylene polymers are known (Moore, P., Polypropylene Handbook, Hanser Publishers Munich 1996, pp. 303-348). Of disadvantage of propylene polymer moldings from common propylene polymers are the insufficient impact properties of the moldings for industrial applications.

- 15 [0003] Known methods for improving the impact properties are the use of blends comprising propylene polymers and butene polymers for producing pipes (EP 0 972 801), the use of blends comprising propylene polymers and hydrogenated butadiene-isoprene block copolymers for blow molded containers (JP 09,227,707) or the use of blends comprising propylene polymers and thermoplastic elastomers such as styrene-ethylene-butene-block copolymers or propylene-ethylene-diene terpolymers for producing injection molded parts (DE 199 27 477). These molded articles have good impact properties, of disadvantage is however the reduced stiffness of the articles.

Object of the invention

- 20 [0004] It is the object of the present invention to provide an industrial polyolefin piping system with improved stiffness, impact strength for use at high service temperature, comprising pipes, fittings, valves and vessels from propylene polymers.

Brief description of the invention

- 25 [0005] According to the present invention, this object is achieved by an industrial polyolefin piping system with improved stiffness, impact strength for use at high service temperature, comprising single- or multilayer pipes, fittings, valves and vessels, consisting of a propylene homopolymer with a melt index of 0.05 to 40 g/10 min at 230°C/2.16 kg, wherein the propylene homopolymers are β -nucleated propylene homopolymers with an $IR\tau > 0.98$ have a tensile modulus ≥ 1500 MPa, a Charpy impact strength at +23°C of ≥ 30 kJ/m² using notched test specimens, a Vicat B temperature $> 90^\circ\text{C}$ and a heat distortion temperature $> 100^\circ\text{C}$.

Detailed description of the invention

- 30 [0006] β -nucleated propylene polymers are isotactic propylene polymers composed of chains in a 3₁ helical conformation having an internal microstructure of β -form spherulites being composed of radial arrays of parallel stacked lamellae. This microstructure can be realized by the addition of β -nucleating agents to the melt and crystallization. The presence of the β -form can be detected through the use of wide angle X-ray diffraction (Moore, J., Polypropylene Handbook, p. 134-135, Hanser Publishers Munich 1996).

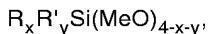
- 35 [0007] The $IR\tau$ of the propylene polymers is measured by Infrared spectroscopy and calculated as described in EP 0 277 514 A2 on page 3.

- 40 [0008] According to a preferred embodiment of the invention the β -nucleated propylene polymers used for producing the industrial piping system have a melt index of 0.05 to 15 g/10min at 230°C/2.16 kg, more preferably 0.1 to 8 g/10 min at 230°C/2.16 kg, most preferably 0.2 to 5 g/10 min at 230°C/2.16 kg.

- 45 [0009] For single- or multilayer pipe fabrication by extrusion propylene homopolymers with a melt index of 0.05 to 40g/10 min at 230°C/2.16 kg, preferably 0.05 to 15 g/min at 230°C/2.16 kg are particularly suitable. For producing fittings and valves by injection molding propylene homopolymers with a melt index of 0.05 to 40 g/10 min at 230°C/2.16 kg are particularly suitable. For vessel fabrication by blow molding a propylene homopolymer with a melt index of 0.05 to 20 g/10 min at 230°C/2.16 kg are particularly suitable.

- 50 [0010] The propylene homopolymers used for producing the industrial piping system show a Charpy impact strength of ≥ 30 kJ/m², preferably ≥ 50 kJ/m², most preferably ≥ 70 kJ/m² at + 23°C.

- 55 [0011] According to an advantageous feature of the present invention, the β -nucleated propylene polymers with an $IR\tau > 0.98$ of the industrial polyolefin piping system are propylene polymers obtained by polymerization with a Ziegler-Natta catalyst system comprising titanium-containing solid components, an organoalumina, magnesium or titanium compound as cocatalyst and an external donor according to the formula



wherein R and R' are identical or different and are branched or cyclic aliphatic or aromatic hydrocarbon residues, and y and x independently from each other are 0 or 1, provided that x + y are 1 or 2.

[0012] A preferred external donor in the Ziegler-Natta catalyst system for producing the propylene polymers for the industrial polyolefin piping system is dicyclopentyldimethoxysilane.

[0013] According to a preferred embodiment of the present invention the β -nucleated propylene polymers of the industrial polyolefin piping system contain 0.01 to 2.0 wt%, based on the polypropylene used,

- dicarboxylic acid derivative type diamide compounds from C₅-C₈-cycloalkyl monoamines or C₆-C₁₂-aromatic monoamines and C₅-C₈-aliphatic, C₅-C₈-cycloaliphatic or C₆-C₁₂-aromatic dicarboxylic acids, and/or
- diamine derivative type diamide compounds from C₅-C₈-cycloalkyl monocarboxylic acids or C₆-C₁₂-aromatic monocarboxylic acids and C₅-C₈-cycloaliphatic or C₆-C₁₂-aromatic diamines, and/or
- amino acid derivative type diamide compounds from amidation reaction of C₅-C₈-alkyl, C₅-C₈-cycloalkyl- or C₆-C₁₂-aryl amino acids, C₅-C₈-alkyl-, C₅-C₈-cycloalkyl- or C₆-C₁₂-aromatic monocarboxylic acid chlorides and C₅-C₈-alkyl-, C₅-C₈-cycloalkyl- or C₆-C₁₂-aromatic mono-amines, as β -nucleating agent.

[0014] Examples of the dicarboxylic acid derivative type diamide compounds from C₅-C₈-cycloalkyl monoamines or C₆-C₁₂-aromatic monoamines and C₅-C₈-aliphatic, C₅-C₈-cycloaliphatic or C₆-C₁₂-aromatic dicarboxylic acids, optionally contained in the β -nucleated propylene polymers of the industrial polyolefin piping system, are

- N,N'-di-C₅-C₈-cycloalkyl-2,6-naphthalene dicarboxamide compounds such as N,N'-dicyclohexyl-2,6-naphthalene dicarboxamide and N,N'-dicyclooctyl-2,6-naphthalene dicarboxamide,
- N,N'-di-C₅-C₈-cycloalkyl-4,4-biphenyldicarboxamide compounds such as N,N'-dicyclohexyl-4,4-biphenyldicarboxamide and N,N'-dicyclopentyl-4,4-biphenyldicarboxamide,
- N,N'-di-C₅-C₈-cycloalkyl-terephthalamide compounds such as N,N'-dicyclohexylterephthalamide and N,N'-dicyclopentylterephthalamide,
- N,N'-di-C₅-C₈-cycloalkyl-1,4-cyclohexanedicarboxamide compounds such as N,N'-dicyclo-hexyl-1,4-cyclohexanedicarboxamide and N,N'-dicyclohexyl-1,4-cyclopentanedicarboxamide.

[0015] Examples of the diamine derivative type diamide compounds from C₅-C₈-cycloalkyl-mono-carboxylic acids or C₆-C₁₂-aromatic monocarboxylic acids and C₅-C₈-cycloaliphatic or C₆-C₁₂-aromatic diamines, optionally contained in the industrial polyolefin piping system, are

- N,N'-C₆-C₁₂-arylene-bis-benzamide compounds such as N,N'-p-phenylene-bis-benzamide and N,N'-1,5-naphthylene-bis-benzamide,
- N,N'-C₅-C₈-cycloalkyl-bis-benzamide compounds such as N,N'-1,4-cyclopentane-bis-benzamide and N,N'-1,4-cyclohexane-bis-benzamide,
- N,N'-p-C₆-C₁₂-arylene-bis-C₅-C₈-cycloalkylcarboxamide compounds such as N,N'-1,5-naphthalene-bis-cyclohexanecarboxamide and N,N'-1,4-phenylene-bis-cyclohexanecarboxamide, and
- N,N'-C₅-C₈-cycloalkyl-bis-cyclohexanecarboxamide compounds such as N,N'-1,4-cyclopentane-bis-cyclohexanecarboxamide and N,N'-1,4-cyclohexane-bis-cyclohexanecarboxamide.

[0016] Examples of the amino acid derivative type diamide compounds, optionally contained in the β -nucleated propylene polymers of the industrial polyolefin piping system, are N-phenyl-5-(N-benzoylamino)pentaneamide and/or

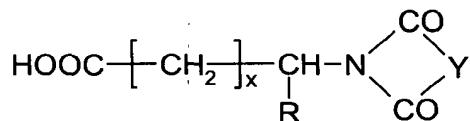
N-cyclohexyl-4-(N-cyclohexyl-carbonylamino)benzamide.

[0017] According to a further advantageous embodiment of the present invention the β -nucleated propylene polymers of the industrial polyolefin piping system contain 0.0001 to 2.0 wt%, based on the polypropylene used, quinacridone type compounds, preferably quinacridone, dimethylquinacridone and/or dimethoxyquinacridone; quinacridonequinone type compounds, preferably quinacridonequinone, a mixed crystal of 5,12-dihydro(2,3b)acridine-7,14-dione with quino(2,3b)acridine-6,7,13,14-(5H,12H)-tetrone as disclosed in EP-B 0 177 961 and/or dimethoxyquinacridonequinone; and/or dihydroquinacridone type compounds, preferably dihydroquinacridone, dimethoxydihydroquinacridone and/or dibenzodihydroquinacridone, as β -nucleating agent.

[0018] According to a further advantageous embodiment of the present invention the β -nucleated propylene polymers of the industrial polyolefin piping system contain 0.01 to 2.0 wt%, based on the polypropylene used, dicarboxylic acid salts of metals from group IIa of periodic system, preferably pimelic acid calcium salt and/or suberic acid calcium salt; and/or mixtures of dicarboxylic acids and salts of metals from group IIa of periodic system, as β -nucleating agent.

- 5 According to a further advantageous embodiment of the present invention the β -nucleated propylene polymers of the industrial polyolefin piping system contain 0.01 to 2.0 wt%, based on the polypropylene used, salts of metals from group IIa of periodic system and imido acids of the formula

10



15

wherein $x = 1$ to 4; $R = H$, -COOH, C_1 - C_{12} -alkyl, C_5 - C_8 -cycloalkyl or C_6 - C_{12} -aryl, and $Y = C_1$ - C_{12} -alkyl, C_5 - C_8 -cycloalkyl or C_6 - C_{12} -aryl - substituted bivalent C_6 - C_{12} -aromatic residues, preferably calcium salts of phthaloylglycine, hexahydrophthaloylglycine, N-phthaloylalanine and/or N-4-methylphthaloylglycine, as β -nucleating agent.

- 20 **[0019]** Preferably the β -nucleated propylene polymers of the industrial piping system are propylene polymers produced by melt mixing propylene homopolymers with 0,0001 to 2,0 % by weight, based on the polypropylenes used, β -nucleating agents at temperatures from 175 to 250°C.

- 25 **[0020]** The propylene polymers used for the inventive industrial polyolefin piping system articles may contain usual auxiliary materials, e.g. 0.01 to 2.5 wt% stabilizers and/or 0.01 to 1 wt% processing aids and/or 0.1 to 1 wt% antistatic agents and/or 0.2 to 3 wt% pigments, in each case based on the propylene polymers used.

- 30 **[0021]** As stabilizers preferably mixtures of 0.01 to 0.6 wt% of phenolic antioxidants, 0.01 to 0.6 wt% of 3-arylbenzofuranones, 0.01 to 0.6 wt% of processing stabilizers based on phosphites, 0.01 to 0.6 wt% of high temperature stabilizers based on disulfides and thioethers and/or 0.01 to 0.8 wt% of sterically hindered amines (HALS) are suitable.

- 35 **[0022]** A further object of the invention is a process for producing an industrial polyolefin piping system with improved stiffness, impact strength for use at high service temperature, comprising single- or multilayer pipe fabrication by extrusion of a propylene homopolymer with a melt index of 0.05 to 40 g/10 min at 230°C/2.16 kg at a melt temperature of 195 to 250°C, fitting and valve fabrication by injection molding of a propylene homopolymer with a melt index of 0.05 to 40 g/10 min at 230°C/2.16 kg at a melt temperature of 220 to 290°C, and vessel fabrication by blow molding of a propylene homopolymer with a melt index of 0.05 to 20 g/10 min at 230°C/2.16 kg at a melt temperature of 185 to 230°C, characterized in that the propylene homopolymers used are β -nucleated propylene homopolymers with an IRt >0.98 having a tensile modulus \geq 1500 MPa, a Charpy impact strength at +23°C \geq 30 kJ/m² using notched test specimens, a Vicat B temperature >90°C and a heat distortion temperature >100°C.

- 40 **[0023]** In the inventive production of industrial polyolefin piping system articles, the extruders for producing the polyolefin pipes can be single screw extruders with an L/D of 20 to 40 or twin screw extruders or extruder cascades of homogenizing extruders (single screw or twin screw). Optionally, a melt pump and/or a static mixer can be used additionally between the extruder and the ring die head. Ring shaped dies with diameters ranging from approximately 16 to 2000 mm and even grater are possible. Advantageous die temperatures for discharging the melt are 180 to 240°C. After leaving the ring-shaped die, the pipes are taken off over a calibrating sleeve and cooled.

- 45 **[0024]** Preferably, for the production of injection molded parts of the inventive process for producing an industrial polyolefin piping system, injection molding machines with injection zones are used, which have three-zone screws with a screw length of 18 to 24 D. Preferred mass temperatures for the melt are 240 to 270°C and temperatures for the injection molds are 20 to 50°C.

- 50 **[0025]** Suitable methods for producing the blow-molded articles of the industrial polyolefin piping system are extrusion blow molding, extrusion stretch blow molding, injection blow molding and injection stretch blow molding. The polyolefin mixtures are extruded through annular dies as a tubular parison, molded into a hollow object by being blown into a divided blowing mold, kept at a temperature of 10 to 55°C and optionally to an additional longitudinal stretching by a stretching stamp and to further radial stretching by blown air. In a second variation, the mixture is injected into an injection mold, heated to 20 to 130°C to produce the parison, and, after being removed and optionally preferably separate heating of the parison in a conditioning mold to temperatures of 80 to 160°C, especially with a first premolding by blowing, transferred into a blowing mold, and by blowing molded into the hollow body, optionally preferably with additional longitudinal stretching by a stretching stamp.

- 55 **[0026]** Preferred applications of industrial polyolefin piping system are chemical plant constructions comprising single- or multilayer pipes, fittings, valves, chambers and vessels with improved stiffness, impact strength and high service temperature for conveyance of fluids.

[0027] The special benefits of the construction parts of the inventive industrial polyolefin piping system are tailor made parts depending on the requirements of service temperatures, conveyed chemical substances and safety aspects.

5 Examples

[0028] The following tests were made using injection molded test specimen prepared according to ISO 1873 Tensile modulus according to ISO 527 (cross head speed 1 mm/min) at +23 °C Charpy impact strength, using notched test specimens according to ISO 179/1eA at +23°C
 10 Vicat B temperature according to ISO 306
 Heat distortion temperature according to ISO 75 Method B

Example 1

15 1.1 Preparation of the β-nucleated propylene polymer

[0029] A mixture of
 94 wt% of a propylene homopolymer, obtained by bulk polymerization using a Ziegler-Natta catalyst system with dicyclopentyldimethoxysilane as external donor, having an IR_t of 0.985, and a melt index of 0.2 g/10 min at 230°C/2.16 kg,
 20 6 wt% of a master batch comprising 98.8 parts by weight of a propylene block copolymer having an ethylene content of 8.3 % by weight, an IR_t of the propylene homopolymer block of 0.985, and a melt index of 0.30 g/10 min at 230°C/2.16 kg, and 0.2 parts by weight of a mixed crystal of 5,12-dihydro(2,3b)acridine-7,14-dione with quino(2,3b)acridine-6,7,13,14-(5H,12H)-tetrone, and
 25 0.05 wt% calcium stearate, 0.1 wt% tetrakis[methylene(3,5-di-t-butylhydroxyhydrocinnamate)] methane and 0.1 wt% tris-(2,4-di-t-butyl-phenyl)-phosphite, based on the sum of the propylene polymers used, is melted in a twin screw extruder with a temperature profile of 100/145/190/215/225/230/230/215/205/190°C, homogenized, discharged and pelletized.

[0030] The resulting polypropylene polymer has a melt index of 0.22 g/10 min at 230°C/2.16 kg, a tensile modulus of 1600 MPa and a Charpy impact strength, using notched test specimens, of 100 kJ/m² at +23°C.

30 1.2 Manufacture of the polyolefin pipe for the industrial polyolefin piping system

[0031] For producing the propylene polymer pipe for the industrial polyolefin piping system, the β-nucleated propylene polymer of 1.1 is introduced in a single screw extruder (L/D=30, D=70mm, temperature profile 200/210/220/220/220/200°C, 40 rpm), melted, extruded through a ring shaped die with a diameter of 110 mm, taken off over a vacuum calibrating sleeve as a pipe of a diameter of 110 mm and a wall thickness of 10 mm, and cooled in a 6 m water bath at 20°C, the taking off velocity being 0.5 m/min.

[0032] Milled test pieces have a Vicat B temperature of 92°C and a Heat Distortion Temperature of 110°C.

40 Example 2

2.1 Preparation of the β-nucleated propylene polymer

[0033] A mixture of
 45 94 wt% of a propylene homopolymer, obtained by bulk polymerization using a Ziegler-Natta catalyst system with dicyclopentyldimethoxysilane as external donor, having an IR_t of 0.985, and a melt index of 8.0 g/10 min at 230°C/2.16 kg, 6 wt% of a master batch comprising 98.8 parts by weight of a propylene homopolymer having an IR_t of 0.987, a melt index of 8.0 g/10 min at 230°C/2.16 kg, and 0.2 parts by weight of a mixed crystal of 5,12-dihydro(2,3b)acridine-7,14-dione with quino(2,3b)acridine-6,7,13,14-(5H,12H)-tetrone and 0.05 wt% calcium stearate, 0.1 wt% tetrakis[methylene(3,5-di-t-butylhydroxyhydrocinnamate)] methane and 0.1 wt% tris-(2,4-di-t-butyl-phenyl)-phosphite, based on the sum of the propylene polymers used, is melted in a twin screw extruder with a temperature profile of 100/145/190/215/225/230/230/215/205/190°C, homogenized, discharged and pelletized.

[0034] The resulting polypropylene polymer has a melt index of 8.2 g/10 min at 230°C/2.16 kg, a tensile modulus of 1800 MPa and a Charpy impact strength, notched, of 88 kJ/m² at +23°C.

55 2.2 Manufacture of the injection molded polyolefin pipe fitting for the industrial polyolefin piping system

[0035] The β-nucleated propylene polymer as described in 2.1 is processed in a injection molding machine, which

has a three-zone screw with a screw length of 22 D, at a mass temperature of 220°C and a mold temperature of 50°C into a pipe fitting having a diameter of 60 mm, a wall thickness of 4 mm and a length of 92 mm.

[0036] Milled test pieces have a Vicat B temperature of 91°C and a heat distortion temperature of 102°C.

5 Example 3

3.1 Preparation of the β -nucleated propylene polymer

[0037] A mixture of

10 75 wt% of a propylene homopolymer obtained by bulk polymerization using a Ziegler-Natta catalyst system with dicyclopentyldimethoxysilane as external donor, having an IR τ of 0.985 and a melt index of 2.8 g/10 min at 230°C/2.16 kg, 25 wt% of a master batch comprising 99.5 parts by weight of a propylene homopolymer obtained by bulk polymerization using a Ziegler-Natta catalyst system with dicyclopentyldimethoxysilane as external donor, having an IR τ of 0.987 and a melt index of 13 g/10 min at 230°C/2.16 kg, and 0.5 parts by weight of hexahydrophthaloylglycine calcium salt, and 15 0.1 wt% calcium stearate, 0.1 wt% tetrakis[methylene(3,5-di-t-butylhydroxyhydrocinnamate)]methane and 0.1 wt% tris-(2,4-di-t-butylphenyl)phosphite, based on the sum of the propylene polymers used, is melted in a twin screw extruder with a temperature profile of 100/145/185/210/220/225/225/200/185°C, homogenized, discharged and pelletized.

20 [0038] The resulting polypropylene polymer has a melt index of 3.8 g/10 min at 230°C/2.16 kg, a tensile modulus of 1700 MPa and a Charpy impact strength, using notched test specimens, of 50 kJ/m² at +23°C.

3.2 Manufacture of the test container with rectangular base for the industrial polyolefin piping system

[0039] An injection stretch blowing equipment comprising a plasticizing unit with a three-zone screw, a turning table with a quadruple injection mold, a conditioning mold with three heating zones, a blowing mold with a stretching stamp and ejection equipment, the β -nucleated propylene polymer of 3.1 is melted in the plasticizing unit at a temperature profile of 100/150/200/225/215°C and injected into the quadruple injection mold, which had been heated to 110°C. The parison, weighing 30 g and having a wall thickness of 4.6 to 5.5 mm and a height of 96 mm, is taken by the sleeve of the turning table from the injection mold and brought by a swiveling motion of 90° of the turning table into the electrically heated conditioning mold, the heating zones of which are adjusted to a temperature of 131°C (bottom of the container parison), of 134°C (center part of the container parison) and of 131°C (upper part of the container parison). After a conditioning period of 85 seconds, preblow-molding by compressed air is followed by removal of the blow-molded preform and transfer by means of the sleeve of the turning table by the swiveling motion of the turning table through 90° into the blowing mold, where the blow-molded preform initially is subjected to longitudinal stretching by the stretching stamp and subsequently molded by compressed air at a pressure of 20 bar. The longitudinal stretching of the preform in the blowing mold is 2.8 : 1 and the radial stretching of the preform is 2.0 : 1. After 8.0 seconds, the blow-molded control container with rectangular base is removed from the open blowing mold, supplied by the swiveling motion of the turning table through 90° to the ejector and ejected.

40 [0040] To determine Vicat B temperature, test pieces were milled after the container shell was cut open. Vicat test results in a Vicat B temperature of 93°C. A heat distortion temperature of 104°C was determined.

Example 4

4.1 Preparation of the β -nucleated propylene polymer

[0041] A mixture of

45 95 wt% of a propylene homopolymer, obtained by bulk polymerization using a Ziegler-Natta catalyst system with dicyclopentyldimethoxysilane as external donor, having an IR τ of 0.986 and a melt index of 0.2 g/10 min at 230°C/2.16 kg, 5 wt% of a master batch comprising 97.5 parts by weight of a propylene homopolymer having an IR τ of 0.987 and a melt index of 4.2g/10 min at 230°C/2.16 kg , and 2.5 parts by weight of N,N'-dicyclohexyl-2,6-naphthalenedicarboxamide, and 0.05 wt% calcium stearate, 0.1 wt% tetrakis[methylene(3,5-di-t-butylhydroxyhydrocinnamate)]methane and 0.1 wt% tris-(2,4-di-t-butylphenyl)-phosphite, based on the sum of the propylene polymers used, is melted in a twin screw extruder with a temperature profile of 100/145/190/215/225/225/205/190°C, homogenized, discharged and pelletized.

55 [0042] The resulting polypropylene polymer has a melt index of 0.28 g/10 min at 230°C/2.16 kg, a tensile modulus of 1750 MPa and a Charpy impact strength, using notched tes specimens, of 90 kJ/m².

4.2 Manufacture of the polyolefin pipe for the industrial polyolefin piping system

[0043] For producing the propylene polymer pipe for the industrial polyolefin piping system, the β -nucleated propylene polymer of 4.1 is introduced in a single screw extruder (L/D=30, D=70mm, temperature profile 200/210/220/220/220/200°C, 40 rpm), melted, extruded through a ring shaped die with a diameter of 110 mm, taken off over a vacuum calibrating sleeve as a pipe of a diameter of 110 mm and a wall thickness of 10 mm, and cooled in a 6 m water bath at 20°C, the taking off velocity being 0.5 m/min.

[0044] Vicat B temperature was 95°C and Heat Distortion Temperature was 114°C.

10

Claims

1. Industrial polyolefin piping system with improved stiffness, impact strength for use at high service temperature, comprising single- or multilayer pipes, fittings, valves, chambers and vessels, consisting of a propylene homopolymer with a melt index of 0.05 to 40 g/10 min at 230°C/2.16 kg, wherein the propylene homopolymers are β -nucleated propylene homopolymers with an IRt >0.98 have a tensile modulus \geq 1500 MPa, a Charpy impact strength at +23°C of \geq 30 kJ/m², using notched test specimens, a Vicat B temperature \geq 90°C and a heat distortion temperature $>$ 100°C.
2. Industrial polyolefin piping system according to claim 1, comprising a propylene homopolymer with a melt index of 0.05 to 15 g/10 min at 230°C/2.16 kg.
3. Industrial polyolefin piping system according to one of the claims 1 or 2, wherein the β -nucleated propylene polymers with an IRt >0.98 are propylene polymers obtained by polymerization with a Ziegler-Natta catalyst system comprising titanium-containing solid components, an organoaluminum, magnesium or titanium compound as co-catalyst and an external donor according to the formula



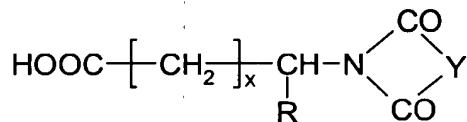
30

wherein R and R' are identical or different and are branched or cyclic aliphatic or aromatic hydrocarbon residues, and y and x independently from each other are 0 or 1, provided that x + y are 1 or 2.

4. Industrial polyolefin piping system according to claim 3, wherein the external donor is dicyclopentylmethoxysilane.
5. Industrial polyolefin piping system according to one of claims 1 to 4, wherein the β -nucleated propylene polymer contains 0.0001 to 2.0 wt%, based on the polypropylene used,
 - dicarboxylic acid derivative type diamide compounds from C₅-C₈-cycloalkyl monoamines or C₆-C₁₂-aromatic monoamines and C₅-C₈-aliphatic, C₅-C₈-cycloaliphatic or C₆-C₁₂-aromatic dicarboxylic acids, and/or
 - diamine derivative type diamide compounds from C₅-C₈-cycloalkyl monocarboxylic acids or C₆-C₁₂-aromatic monocarboxylic acids and C₅-C₈-cycloaliphatic or C₆-C₁₂-aromatic diamines, and/or
 - amino acid derivative type diamide compounds from amidation reaction of C₅-C₈-alkyl-, C₅-C₈-cycloalkyl- or C₆-C₁₂-arylamino acids, C₅-C₈-alkyl-, C₅-C₈-cycloalkyl- or C₆-C₁₂-aromatic monocarboxylic acid chlorides and C₅-C₈-alkyl-, C₅-C₈-cycloalkyl- or C₆-C₁₂-aromatic monoamines, as β -nucleating agent.
6. Industrial polyolefin piping system according to one of the claims 1 to 4, wherein in that the β -nucleated propylene polymer contains 0.0001 to 2.0 wt%, based on the polypropylene used, quinacridone type compounds, preferably quinacridone, dimethylquinacridone and/or dimethoxyquinacridone; quinacridonequinone type compounds, preferably quinacridonequinone, a mixed crystal of 5,12-dihydro(2,3b)acridine-7,14-dione with quino(2,3b)acridine-6,7,13,14-(5H,12H)-tetrone and/or dimethoxyquinacridonequinone; and/or dihydroquinacridone type compounds, preferably dihydroquinacridone, dimethoxydihydroquinacridone and/or dibenzodihydroquinacridone, as β -nucleating agent.
7. Industrial polyolefin piping system according to one of the claims 1 to 4, wherein the β -nucleated propylene polymer contains 0.01 to 2.0 wt%, based on the polypropylene used, dicarboxylic acid salts of metals from group IIa of periodic system, preferably pimelic acid calcium salt and/or suberic acid calcium salt; and/or mixtures of dicarboxylic acids and salts of metals from group IIa of periodic system, as β -nucleating agent.

8. Industrial polyolefin piping system according to one of the claims 1 to 4, characterized in that the β -nucleated propylene polymer contains 0.01 to 2.0 wt%, based on the polypropylene used, salts of metals from group IIa of periodic system and imido acids of the formula

5



10

wherein $x = 1$ to 4; R = H, -COOH, C₁-C₁₂-alkyl, C₅-C₈-cycloalkyl or C₆-C₁₂-aryl, and Y = C₁-C₁₂-alkyl, C₅-C₈-cycloalkyl or C₆-C₁₂-aryl - substituted bivalent C₆-C₁₂-aromatic residues, preferably calcium salts of phthaloylglycine, hexahydrophthaloylglycine, N-phthaloylalanine and/or N-4-methylphthaloylglycine, as β -nucleating agent.

15

9. Process for producing an industrial polyolefin piping system with improved stiffness, impact strength for use at high service temperature, comprising single- or multilayer pipe fabrication by extrusion of a propylene homopolymer with a melt index of 0.05 to 40 g/10 min at 230°C/2.16 kg at a melt temperature of 195 to 250°C, fitting and valve fabrication by injection molding of a propylene homopolymer with a melt index of 2 to 40 g/10 min at 230°C/2.16 kg at a melt temperature of 220 to 290°C, and vessel fabrication by blow molding of a propylene homopolymer with a melt index of 1 to 20 g/10 min at 230°C/2.16 kg at a melt temperature of 185 to 230°C, characterized in that the propylene homopolymers used are β -nucleated propylene homopolymers with an IRt > 0.98 having a tensile modulus \geq 1500 MPa, a Charpy impact strength, using notched test specimens, at 23°C \geq 30 kJ/m², a Vicat B temperature > 90°C and a heat distortion temperature > 100°C.
10. Use of an industrial polyolefin piping system according to one of claims 1 to 7 for chemical plant constructions comprising pipes, fittings, valves and vessels with improved stiffness, impact strength and high service temperature for conveyance of fluids

20

25

30

35

40

45

50

55



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 01 11 2365

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	WO 99 40151 A (BERNREITNER KLAUS ;KONRAD ROLAND (AT); EBNER KARL (AT); PCD POLYME) 12 August 1999 (1999-08-12) * page 2, line 20-22; claims 1,2; example 1 *	1-5,9,10	C08K5/00 C08L23/10 F16L9/12 F16L9/127 F16L9/133 B32B1/08 B32B27/32 C08F110/06 C08F297/08 C08F10/06
X	WO 99 24479 A (BOREALIS AS ;JAEAESKELAEINEN PIRJO (FI); MALM BO (FI); HUOVINEN PA) 20 May 1999 (1999-05-20) * claims 1,3,13-15,22,24,25,31,32,36-41; examples 9,10,13; tables III,V,VI *	1-4,9,10	
A	EP 0 714 923 A (SHOWA DENKO KK) 5 June 1996 (1996-06-05) * page 7, line 48 - page 8, line 5; claims; examples; tables *	1-3,7,9, 10	
A	DE 297 22 949 U (PCD POLYMER GMBH) 26 February 1998 (1998-02-26) * page 2, paragraph 2 - page 3, paragraphs 3-5; claims 1-3,5; table 2 *	1,2,9,10	
TECHNICAL FIELDS SEARCHED (Int.Cl.7)			
C08K C08L F16L B32B C08F			
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	31 October 2001	Derz, T	
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone	T : theory or principle underlying the invention		
Y : particularly relevant if combined with another document of the same category	E : earlier patent document, but published on, or after the filing date		
A : technological background	D : document cited in the application		
O : non-written disclosure	L : document cited for other reasons		
P : intermediate document	& : member of the same patent family, corresponding document		

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 01 11 2365

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

31-10-2001

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 9940151	A	12-08-1999	WO	9940151 A1	12-08-1999
			EP	1044240 A1	18-10-2000
			NO	20003821 A	26-07-2000

WO 9924479	A	20-05-1999	AU	1035799 A	31-05-1999
			BR	9814854 A	03-10-2000
			CN	1285851 T	28-02-2001
			EP	1028985 A1	23-08-2000
			WO	9924479 A1	20-05-1999

EP 0714923	A	05-06-1996	AU	705455 B2	20-05-1999
			AU	3790395 A	30-05-1996
			CA	2162946 A1	22-05-1996
			CN	1128767 A	14-08-1996
			DE	69517771 D1	10-08-2000
			DE	69517771 T2	23-11-2000
			EP	0714923 A1	05-06-1996
			JP	3002119 B2	24-01-2000
			JP	8198922 A	06-08-1996
			US	5703172 A	30-12-1997
			US	5804665 A	08-09-1998
			JP	3034452 B2	17-04-2000
			JP	8198923 A	06-08-1996

DE 29722949	U	26-02-1998	DE	29722949 U1	26-02-1998

PUB-NO: EP001260545A1
DOCUMENT-IDENTIFIER: EP 1260545 A1
TITLE: Industrial polyolefin piping system
PUBN-DATE: November 27, 2002

ASSIGNEE-INFORMATION:

NAME	COUNTRY
BOREALIS TECH OY	FI

APPL-NO: EP01112365
APPL-DATE: May 21, 2001

PRIORITY-DATA: EP01112365A (May 21, 2001)

INT-CL (IPC): C08K005/00 , C08L023/10 ,
F16L009/12 , F16L009/127 ,
F16L009/133 , B32B001/08 ,
B32B027/32 , C08F110/06 ,
C08F297/08 , C08F010/06

EUR-CL (EPC): B32B001/08 , B32B027/32 ,
C08F004/646 , C08F110/06 ,
C08F110/06 , C08K005/00 ,
F16L009/12 , F16L009/127 ,
F16L009/133

ABSTRACT:

CHG DATE=20030114 STATUS=O> Industrial polyolefin piping system with improved stiffness,

impact strength for use at high service temperature, comprising single- or multilayer pipes, fittings, chambers, valves and vessels, consisting of beta -nucleated propylene homopolymers with an IR tau >0.98 having a tensile modulus ≥ 1500 MPa, a Charpy impact strength, notched, at +23 DEG C ≥ 30 kJ/m² and a Vicat B temperature >90 DEG C and a Heat Distortion Temperature >100 DEG C. The industrial polyolefin piping system is suitable for chemical plant constructions comprising single- or multilayer pipes, fittings, chambers, valves and vessels with improved stiffness, impact strength and high service temperature, preferred for conveyance of natural gas, dangerous liquids and/or toxic liquids.